

## **Case 8056**

### **Haemorrhagic cellular leiomyoma**

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**Section:** Genital (Female) Imaging

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**Patient:** 27 year(s), female

## **Clinical History**

A 27-year-old Caucasian female patient presenting with a large uterine mass on gynecological ultrasound, which showed significant growth in relation to the previous examination.

## **Imaging Findings**

The patient was asymptomatic. She was not taking oral contraceptives and was not pregnant or in postpartum period. Physical examination revealed a non tender pelvic mass. Blood tests were normal.

The annual follow-up gynaecologic ultrasound examination of a previously documented uterine leiomyoma revealed a significantly larger uterine mass, slightly hyper-echoic compared to the myometrium. The patient was then referred for an MR study, which showed an intramural lesion with regular margins, hypo-intense on T1-weighted images (Fig. 2), hyper-intense on T2-weighted images (Fig. 3), enhancing intensively with gadolinium (Fig. 4).

After treatment with GnRH analogues, which induced tumor reduction, a myomectomy was performed (Figs. 5 and 6). The isolated solid tumor measured 8.5 cm and was well circumscribed, with areas of haemorrhage (20% of total). Histologically, smooth muscle cells showed nuclear atypia and mitoses in number of 2/10 x 400 fields. Areas of haemorrhage were not associated with necrosis.

Based on the histological pattern of the tumor, patient age and response to treatment, the final diagnosis was haemorrhagic cellular leiomyoma.

## Discussion

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Leiomyomas are the most common gynecologic neoplasm, occurring in 20-30% of women in reproductive age. They are predominantly composed of smooth muscle cells and variable amounts of fibrous connective tissue [1, 2].

Haemorrhagic cellular leiomyomas are a histological subtype of leiomyomas. They occur almost exclusively in young women treated with oral contraceptives, during pregnancy or in postpartum period (these features were not present in our case). They are referred as apoplectic leiomyomas" due to the fact that the clinical symptoms are often acute and pathological features are dominated by recent haemorrhages. These lesions are different from extensive or zonal infarcts, commonly termed red degeneration", which coexist with coagulative or ischemic necrosis, with no hypercellularity [3, 4].

Typically, non-degenerated uterine leiomyomas appear on MR imaging as well-circumscribed masses of homogeneously decreased signal intensity compared to that in the outer myometrium on T2-weighted images and of intermediate signal on T1-weighted images. Cellular leiomyomas, which are composed of compact smooth muscle cells with little or no collagen, can have relatively higher intensity signal on T2-weighted images and demonstrate enhancement on contrast enhanced images [1].

MR imaging can help differentiate cellular leiomyomas from degenerated T2-weighted hyper-intense leiomyomas, as degenerated leiomyomas often show irregular, peripheral, or minimal enhancement compared with that of myometrium, depending on the degeneration within the tumor [5].

The differential diagnosis also includes lesions with extensive oedema affecting their signal intensity, which can be high on T2 weighted images and demonstrate marked enhancement. Microscopy will show fluid in the stroma of the leiomyoma, often in association with collagen [2].

Finally, although it has been suggested that an irregular margin of a uterine leiomyoma at MR imaging is related to a sarcomatous transformation, the ability of MR imaging to allow differentiation of cellular (or degenerated) leiomyoma from leiomyosarcoma of the uterus has not been yet assessed. The diagnosis is established histologically [1, 5].

When haemorrhage and hypercellularity coexist, the suspicion of leiomyosarcoma is increased. The major differences between cellular haemorrhagic leiomyomas and leiomyosarcomas are that haemorrhagic leiomyomas have smaller cells, less atypia, fewer abnormal mitotic figures, small size, multiple arrangement, multiple discrete areas of haemorrhage and discrete periphery. Also, their mitotic activity is confined to a narrow zone adjacent to the haemorrhage [3].

## Final Diagnosis

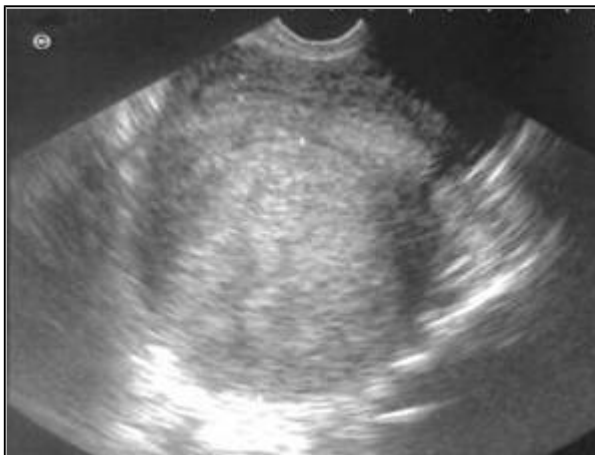
Haemorrhagic cellular leiomyoma

## Figures

Figure 1 No title

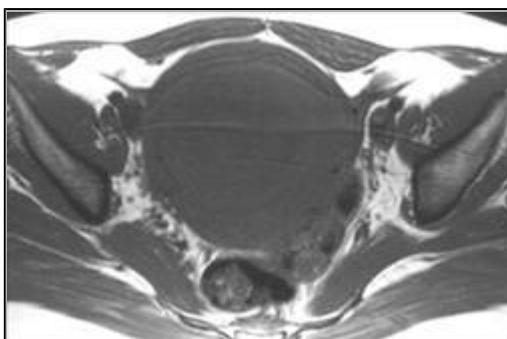


Ultrasound: sagittal trans-abdominal (a) and trans-vaginal (b), showing a large posterior uterine mass, slightly hyperechoic compared to the myometrium.



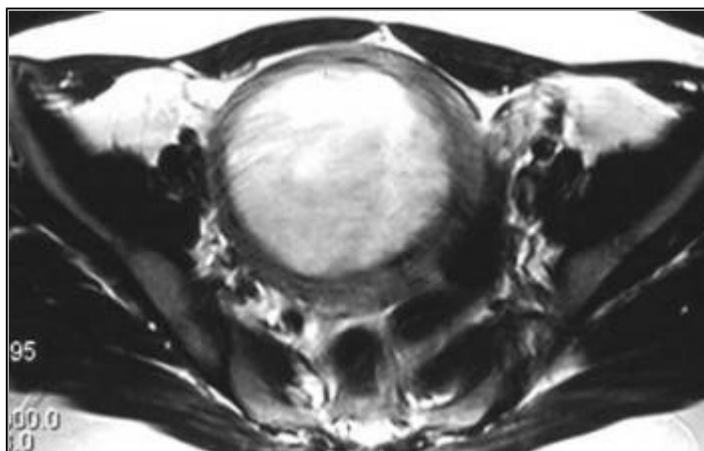
Ultrasound: sagittal trans-abdominal (a) and trans-vaginal (b), showing a large posterior uterine mass, slightly hyperechoic compared to the myometrium.

Figure 2 No title



MR T1-weighted axial image shows a large uterine mass with low signal intensity.

**Figure 3 No title**

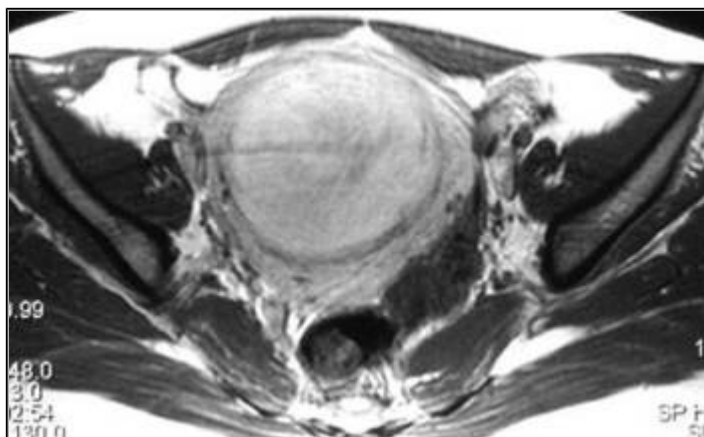


MR T2-weighted axial (a) and sagittal (b) images show a well circumscribed, hyperintense heterogeneous posterior intramural uterine mass.



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**Figure 4 No title**



Contrast-enhanced T1-weighted image shows intense enhancement of the mass.

**Figure 5 No title**



Clinical inspection reveals a visible swelling on lower abdomen.

**Figure 6 No title**



Intra-operative aspect (a). The gross specimen of myomectomy (b) shows dark red softening macroscopic appearance.



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**Uterine Neoplasms [C13.371.852.762]**

Tumors or cancer of the UTERUS.

**References**

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**Citation**

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